

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: June 29, 2002, 22:05:57 ; Search time 17398.4 seconds
(without alignments)
1042.620 Million cell updates/sec

Title: US-09-303-518D-127

Perfect score: 1344

Sequence: 1 atgattaaatcaaaaagg.....ccnttgagaaggagctga 1344

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database :

EST:*

1: em_estba:*

2: em_esthum:*

3: em_estin:*

4: em_estmu:*

5: em_estov:*

6: em_estpl:*

7: em_estro:*

8: em_hic:*

9: gb_estl:*

10: gb_est2:*

11: gb_hic:*

12: gb_gss:*

13: em_gss_hum:*

14: em_gss_inv:*

15: em_gss_pln:*

16: em_gss_vrt:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	108.6	8.1	349	12	CNS07GYI
2	39.2	2.9	540	10	BF260729 HVSMEF002
3	39.2	2.9	544	10	BF260723 HVSMEF002
4	39.2	2.9	551	10	BF253989 HVSMEF000
5	39.2	2.9	574	10	BF265938 HV_Cea001
6	39.2	2.9	622	10	BI957371 HVSMEF000
7	39.2	2.9	634	10	BI959896 HVSMEF002
8	39.2	2.9	636	10	BI959473 HVSMEF001
9	39.2	2.9	704	10	BF262832 HVSMEF000
10	39.2	2.9	759	10	BF259476 HVSMEF001
11	39.2	2.9	764	10	BF616515 HVSMEC001
12	39.2	2.9	783	10	BF626740 HVSMEB000
13	39.2	2.9	841	10	BF254527 HVSMEF000
14	39.2	2.9	847	10	BI957008 HVSMEF000
15	39.2	2.9	864	10	BI957786 HVSMEF001
16	39.2	2.9	888	9	AW983120 HVSMEB000
17	39	2.9	506	9	AI399239 NCW10F2T3

18	39	2.9	621	10	BM490565
19	38.2	2.8	455	10	BI956450
20	37.6	2.8	489	9	AV939450
21	37.6	2.8	492	10	BE348841
22	37.6	2.8	575	10	BE551347
23	37.6	2.8	697	10	BF628426
24	37.2	2.8	841	10	BG342965
25	37	2.8	430	12	AZ049555
26	37	2.8	497	12	P9478
27	36.8	2.7	893	9	AL523270
28	36.6	2.7	612	12	AO161443
29	36.4	2.7	388	10	BM375129
30	36.2	2.7	682	9	AV917449
31	36	2.7	284	9	BE063171
32	36	2.7	298	9	AA831113
33	36	2.7	358	10	BF109446
34	36	2.7	384	9	AI392198
35	36	2.7	393	9	AI306569
36	36	2.7	400	9	AI609778
37	36	2.7	419	9	AA468664
38	36	2.7	477	9	AI300364
39	36	2.7	486	9	AI039130
40	36	2.7	488	9	AI378387
41	36	2.7	497	9	AI701002
42	36	2.7	526	9	AW006812
43	36	2.7	544	9	AI743812
44	36	2.7	925	12	CNS0091P
45	35.8	2.7	932	12	CNS01KEO
46	35.8	2.7	1013	12	CNS04MSY
47	35.6	2.6	413	9	AV631377
48	35.6	2.6	606	12	AO160470
49	35.6	2.6	695	12	AO162608
50	35.4	2.6	309	10	R23876
51	35.4	2.6	421	10	BE423134
52	35.4	2.6	479	10	BF293599
53	35.4	2.6	526	10	BG262921
54	35.4	2.6	561	10	BG262203
55	35.4	2.6	580	10	BE430910
56	35.4	2.6	599	10	BF483435
57	35.4	2.6	617	12	AZ935303
58	35.4	2.6	642	9	AA263267
59	35.4	2.6	702	10	BE414023
60	35.4	2.6	704	10	BE413893
61	35.4	2.6	849	10	BE642582
62	35.2	2.6	613	10	BG577607
63	35.2	2.6	642	9	AV273615
64	35.2	2.6	684	12	AZ085775
65	35.2	2.6	797	10	BE412725
66	35	2.6	356	10	H83299
67	35	2.6	429	9	AI816996
68	35	2.6	458	9	AW174071
69	35	2.6	620	10	BE425512
70	35	2.6	802	9	AW483111
71	35	2.6	826	10	BF624680
72	35	2.6	892	10	BI836975
73	34.8	2.6	275	9	AW574530
74	34.8	2.6	488	9	AI452688
75	34.6	2.6	323	9	AI368671
76	34.6	2.6	376	10	BF293584
77	34.6	2.6	407	9	AW172028
78	34.6	2.6	433	9	AW238848
79	34.6	2.6	497	10	BE378302
80	34.6	2.6	501	10	BE396602
81	34.6	2.6	525	10	BE407307
82	34.6	2.6	554	10	BE792826
83	34.6	2.6	582	10	BM042302
84	34.6	2.6	605	10	BI195835
85	34.6	2.6	605	10	BE907017
86	34.6	2.6	609	12	FR0041777
87	34.6	2.6	624	10	BE394122
88	34.6	2.6	627	10	BE302953
89	34.6	2.6	636	10	BE255207
90	34.6	2.6	659	10	BE254129

<http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Close TJ, Wing R, Kleinof A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html>)"

BASE COUNT

111 a 165 c 174 g 90 t

Query Match Best Local Similarity 50.0%; Score 39.2; DB 10; Length 540; Matches 98; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

QY 908 cggatattgaagcgcgacatcaacaagcgacacattattggagccctaccacatc 967
 Db 167 CGGTGATCTTCGACGCCGAGACGCGGCGGTGACGCTTCCGTTGACAAAGA 226
 QY 968 agattccgttatgaagaagcgcgacgaagcgtgttcgctggttgcgcgcagc 1027
 Db 227 AGATGAGAGCTCTGACGCTGCGGCGACGACATGAGGTCTTCCCAAGGTGAGCGGGG 286
 QY 1028 cggacaatactccatcacgcgttacgacctgagccattctctgaanaaaccttca 1087
 Db 287 TGAGCTCTGACGCCGAGAGCGCCGAGAGCCGCAAGTCAACACCCCTGCTGAGA 346
 QY 1088 agttacgacagccgt 1103
 Db 347 AGGCCAAGAGCCCGT 362

RESULT 3

BF260723

LOCUS

BF260723 544 bp mRNA linear EST 22-OCT-2001
 HVSMEF0022M09f Hordeum vulgare seedling root EST library HVCNMA0007
 (Etiolated and unstressed) Hordeum vulgare cDNA clone
 HVSMEF0022M09f, mRNA sequence.

ACCESSION

BF260723

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Hordeum vulgare
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae;
 Triticeae; Hordeum.
 1 (bases 1 to 544)
 Wing, R., Close, T.J., Kleinof, A., Wise, R., Begum, D., Frisch, D., Yu,
 Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
 R.D., Oates, R. and Main, D.
 Development of a genetically and physically anchored EST resource
 for barley genomics: Morex unstressed seedling root cDNA library
 Unpublished (2001)
 On Nov 16, 2000 this sequence version replaced gi:11189836.
 Contact: Wing RA
 Clemson University Genomics Institute
 Clemson University
 100 Jordan Hall, Clemson, SC 29634, USA
 Tel: 864 656 7288
 Fax: 864 656 4293
 Email: rwing@clemson.edu
 Total hg bases = 250
 Seq primer: ATTATACCTCCTCAAGG
 High quality sequence stop: 512.
 Location/Qualifiers
 1..544
 /organism="Hordeum vulgare"
 /cultivar="Morex"
 /db_xref="taxon:4513"
 /clone="HVSMEF0022M09f"
 /clone_1b="Hordeum vulgare seedling root EST library
 HVCNMA0007 (Etiolated and unstressed)"
 /tissue_type="Seedling root"
 /lab_host="TTC121"

/note="Vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI.
 Seeds were surface sterilized then germinated under axenic
 conditions in the dark at room temperature on filter paper
 with water, nystatin and cefotaxime in covered
 crystallization dishes. Five-day old seedling roots were
 then harvested, total RNA was prepared, poly(A) RNA was
 purified, one primary unamplified cDNA library was made,
 and 1 million p1u were in vivo excised to give phagescript
 SK(-) cDNA phagemids. These steps were performed in the TJ
 Close Laboratory at the University of California,
 Riverside (Choi, Close, Fenton). Phagemids were plated and
 picked at the Clemson University Genomics Institute (CUGI)
 (Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
 preparations, DNA sequencing and sequence analysis were
 performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
 , Rambo, Main). The sequence has been trimmed to remove
 vector sequence and contains a minimum of 100 bases of
 phred value 20 or above. For more details on library
 preparation and sequence analysis see
<http://www.genome.clemson.edu/projects/barley>. To order
 this clone see <http://www.genome.clemson.edu/orders> Also
 see Close TJ, Wing R, Kleinof A, Wise R (2001)
 Genetically and physically anchored EST resources for
 barley genomics. Barley Genetics Newsletter 31:29-30.
 (<http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html>)"

BASE COUNT

110 a 165 c 174 g 92 t

Query Match Best Local Similarity 50.0%; Score 39.2; DB 10; Length 544; Matches 98; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

QY 908 cggatattgaagcgcgacatcaacaagcgacacattattggagccctaccacatc 967
 Db 167 CGGTGATCTTCGACGCCGAGACGCGGCGGTGACGCTTCCGTTGACAAAGA 226
 QY 968 agattccgttatgaagaagcgcgacgaagcgtgttcgctggttgcgcgcagc 1027
 Db 227 AGATGAGAGCTCTGACGCTGCGGCGACGACATGAGGTCTTCCCAAGGTGAGCGGGG 286
 QY 1028 cggacaatactccatcacgcgttacgacctgagccattctctgaanaaaccttca 1087
 Db 287 TGAGCTCTGACGCCGAGAGCGCCGAGAGCCGCAAGTCAACACCCCTGCTGAGA 346
 QY 1088 agttacgacagccgt 1103
 Db 347 AGGCCAAGAGCCCGT 362

RESULT 4

BF253989

LOCUS

BF253989 551 bp mRNA linear EST 22-OCT-2001
 HVSMEF0022L01f Hordeum vulgare seedling root EST library HVCNMA0007
 (Etiolated and unstressed) Hordeum vulgare cDNA clone
 HVSMEF0022L01f, mRNA sequence.

ACCESSION

BF253989

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Hordeum vulgare
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae;
 Triticeae; Hordeum.
 1 (bases 1 to 551)
 Wing, R., Close, T.J., Kleinof, A., Wise, R., Begum, D., Frisch, D., Yu,
 Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
 R.D., Oates, R. and Main, D.
 Development of a genetically and physically anchored EST resource
 for barley genomics: Morex unstressed seedling root cDNA library
 Unpublished (2001)
 On Nov 16, 2000 this sequence version replaced gi:11183094.
 Contact: Wing RA

Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 300
Seq primer: AATTAACTCTCACTAAAGG
High quality sequence stop: 540.
Location/Qualifiers

FEATURES

1. 551
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMF0002L01f"
/clone.lib="Hordeum vulgare seedling root EST library
HVCNDA0007 (Etiolated and unstressed)"
/tissue_type="Seedling root"
/lab_host="TUC121"
/note="Vector: lambdaZAP; Site.1: EcoRI; Site.2: XhoI;
Seeds were surface sterilized then germinated under aseptic
conditions in the dark at room temperature on filter paper
with water, nystatin and ceftaxime in covered
crystallization dishes. Five-day old seedling roots were
then harvested, total RNA was prepared, poly(A) RNA was
purified, one primary unamplified cDNA library was made,
and 1 million pfu were in vivo excised to give Bluescript
SK(-) cDNA phagemids. These steps were performed in the TU
Close laboratory at the University of California,
Riverside (Choi, Close, Fenton). Phagemids were plated and
picked at the Clemson University Genomics Institute (CUGI)
(Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
preparations, DNA sequencing and sequence analysis were
performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
, Rambo, Main). The sequence has been trimmed to remove
vector sequence and contains a minimum of 100 bases of
phred value 20 or above. For more details on library
preparation and sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders Also
see Close TJ, Wing R, Kleinof A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html)"

BASE COUNT 128 a 151 c 175 g 96 t 1 others
ORIGIN

Query Match 2.9%; Score 39.2; DB 10; Length 551;
Best Local Similarity 50.0%; Pred. No. 4.3;
Matches 98; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

Oy 908 cagatitgaacgagcgatatacacaagcgacgattatggagcgtacacacatc 967
Db 25 CGGTATTTTGACGCGGAGCAGCGGCGACGCTTCCGGTTGACAGAGA 84
Oy 968 agattccgtatgaagaagcgacgacgagctgttcgctgttcgacgacg 1027
Db 85 AGATGAGACTGCTGACGCTGGCGACGACATGAGGTTCGCCAAGTGGAGCCGGGCG 144
Oy 1028 cggaaanaatcatcatcaacgagctacgacccctcgccattcctgaanaaactctca 1087
Db 145 TGAGTCTGTACGCGGAGCGCGGAGGAGCGCCCAAGTCCATACACCCCTGCTGAGA 204
Oy 1088 agtcaacgacagccgt 1103
Db 205 AGGCCAAGAGGCGCGT 220

RESULT 5
BF265938 574 bp mRNA linear EST 23-OCT-2001
LOCUS BF265938
DEFINITION HV_CEA0013L17f Hordeum vulgare seedling green leaf EST library

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

HVCNDA0004 (Blumeria challenged) Hordeum vulgare cDNA clone
HV_CEA0013L17f, mRNA sequence.
BF265938
BF265938.2 GI:13262431
EST.

REFERENCE

Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poideae
; Triticeae; Hordeum.
1 (bases 1 to 574)

TITLE

JOURNAL

COMMENT

On Nov 17, 2000 this sequence version replaced g1:11196932.
Contact: Wing R
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 348
Seq primer: AATTAACTCTCACTAAAGG
High quality sequence stop: 538.
Location/Qualifiers

FEATURES

source

1. 574
/organism="Hordeum vulgare"
/cultivar="C116155 (M1a13)"
/db_xref="taxon:4513"
/clone="HV_CEA0013L17f"
/clone.lib="Hordeum vulgare seedling green leaf EST
library HVCNDA0004 (Blumeria challenged)"
/tissue_type="Seedling green leaf"
/lab_host="TUC121"
/note="Vector: lambdaZAP; Site.1: EcoRI; Site.2: XhoI;
C.I. 16155 (M1a13) plants were greenhouse grown in the R
Wise lab at Iowa State University, Ames, IA; 7 day old
green seedlings were challenged with isolate A27 (AvrM1a13
) of Blumeria graminis f. sp. hordei, and leaves were
harvested 20 and 24 hr post-inoculation and snap frozen;
uninoculated leaves were harvested 20 hr post-inoculation
(Wing, Wise). In the TU Close lab at the University of
California, Riverside, total RNA was prepared from each
sample pool, equal quantities of all three RNA pools were
combined, poly(A) RNA was purified from the mixture, one
cDNA library was made, and 1 million pfu were in vivo
excised to give Bluescript SK(-) cDNA phagemids (Choi,
Close). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main).
The sequence has been trimmed to remove vector sequence
and contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders Also
see Close TJ, Wing R, Kleinof A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html)"

BASE COUNT
ORIGIN

119 a 177 c 178 g 100 t

Query Match 2.9%; Score 39.2; DB 10; Length 574;
Best Local Similarity 50.0%; Pred. No. 4.4;

[illegible]

/lab_host="TJUC121"

/note="Vector: pBluescript SK(-); Site.1: EcoRI; Site.2: XhoI; Plants were grown at Washington State University, Pullman, WA in a greenhouse, the rachises were excised and frozen in liquid nitrogen (Kleinof's lab). In the TJ Close lab at the University of California, Riverside total RNA was prepared, poly(A) was purified, one primary unamplified cDNA library was made, and 1 million pfu were in vivo excised to give pBluescript SK(-) cDNA phagemids (Chin). Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phred value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Close TJ, Wing R, Kleinof's A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html>)"

BASE COUNT 131 a 185 c 205 g 113 t

ORIGIN

Query Match 2.9%; Score 39.2; DB 10; Length 634;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 98; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

QY 908 cggatttaagcggcgattacacaaagcgccgattattggagcgctaccacaac 967
DB 177 CGGTATCTTCGACGCCGGAAGCAGCGGCGCTTCGCTTCGGTTCCAGACA 236
QY 968 agatttcggtatcgaagaagcgccgaagaagctgtcgtggttgccgcagc 1027
DB 237 AGATGGAGCTCGTCCGACCTCGGCGAGCATCGAGGTCTTCCGCAAGGTGAGCGGGC 296
QY 1028 cggacaataactcactacacgcgtacgacctcgccatttcctgaacaaactctca 1087
DB 297 TGAGCTCGTACGCCGAGAGCGCCGCGCAAGTCCATCACCCTCTGTGAGA 356
QY 1088 agttcagcagacgcgt 1103
DB 357 AGGCCAAGAGCGCGCT 372

RESULT 8

LOCUS B1959473 636 bp mRNA linear EST 22-OCT-2001
DEFINITION HVSMEN0019N03f Hordeum vulgare rachis EST library HVCNMA0015 (normal) Hordeum vulgare cDNA clone HVSMEN0019N03f, mRNA sequence.

ACCESSION B1959473
VERSION B1959473.1 GI:16310728

KEYWORDS EST

SOURCE

ORGANISM Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poideae

1 (bases 1 to 636)
; Triticeae; Hordeum.

REFERENCE Wing, R., Close, T.J., Kleinof's, A., Wise, R., Chin, A., Begum, D., Frisch, D., Atkins, M., Yu, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Oates, R. and Main, D.

TITLE Development of a genetically and physically anchored EST resource for barley genomics: Morex rachis cDNA library

JOURNAL Unpublished (2001)

COMMENT Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA

Tel: 864 656 7288
Fax: 864 656 4293
Email: twing@clemson.edu
Total hg bases = 585
Seq primer: AATTACCCCTCCTAAAGCG
High quality sequence start: 5
High quality sequence stop: 624.
Location/Qualifiers
1. 636

FEATURES

source

/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEN0019N03f"
/clone_lib="Hordeum vulgare rachis EST library HVCNMA0015 (normal)"
/tissue_type="Rachis"
/lab_host="TJUC121"

/note="Vector: pBluescript SK(-); Site.1: EcoRI; Site.2: XhoI; Plants were grown at Washington State University, Pullman, WA in a greenhouse, the rachises were excised and frozen in liquid nitrogen (Kleinof's lab). In the TJ Close lab at the University of California, Riverside total RNA was prepared, poly(A) was purified, one primary unamplified cDNA library was made, and 1 million pfu were in vivo excised to give pBluescript SK(-) cDNA phagemids (Chin). Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phred value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Close TJ, Wing R, Kleinof's A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html>)"

BASE COUNT 132 a 188 c 203 g 113 t

ORIGIN

Query Match 2.9%; Score 39.2; DB 10; Length 636;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 98; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

QY 908 cggatttaagcggcgattacacaaagcgccgattattggagcgctaccacaac 967
DB 175 CGGTATCTTCGACGCCGGAAGCAGCGGCGCTTCGCTTCGGTTCCAGACA 234
QY 968 agatttcggtatcgaagaagcgccgaagaagctgtcgtggttgccgcagc 1027
DB 235 AGATGGAGCTCGTCCGACCTCGGCGAGCATCGAGGTCTTCCGCAAGGTGAGCGGGC 294
QY 1028 cggacaataactcactacacgcgtacgacctcgccatttcctgaacaaactctca 1087
DB 295 TGAGCTCGTACGCCGAGAGCGCCGCGCAAGTCCATCACCCTCTGTGAGA 354
QY 1088 agttcagcagacgcgt 1103
DB 355 AGGCCAAGAGCGCGCT 370

RESULT 9

LOCUS BF628632 704 bp mRNA linear EST 22-OCT-2001
DEFINITION HVSMEN0006N12f Hordeum vulgare seedling shoot EST library HVCNMA0002 (Dehydration stress) Hordeum vulgare cDNA clone

ACCESSION BF628632
VERSION BF628632.2 GI:13090280

KEYWORDS EST.
SOURCE barley.
ORGANISM Hordeum vulgare
REFERENCE Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae
AUTHORS ; Triticeae; Hordeum.
1 (bases 1 to 704)
Wing, R., Close, T.J., Kleinbols, A., Wise, R., Begum, D., Frisch, D., Yu
, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
, R.D., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex drought-stressed seedling shoot cDNA
library
JOURNAL Unpublished (2001)
COMMENT On Dec 19, 2000 this sequence version replaced gi:11892790.
Contact: Wing RA
Clemson University Genomics Institute
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 580
Seq primer: AATTAACCTCCTCAAGG
High quality sequence stop: 663.
FEATURES
Source
1..704
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSME0006N12f"
/clone_lib="Hordeum vulgare seedling shoot EST library
HVCNMA0002 (Dehydration stress)"
/tissue_type="Seedling shoot"
/lab_host="TJC121"
/note="Vector: lambdaZAP; Site 1: EcoRI; Site 2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedlings were
incubated at 90% RH for 24 hr. Shoots were then harvested,
total RNA was prepared, poly(A) RNA was purified, one
primary unamplified cDNA library was made, 600000 pfu were
in vivo excised to give plusescript SK(-) cDNA phagemids.
These steps were performed in the TJ Close laboratory at
the University of California, Riverside (Choi, Close,
Fenton). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main).
The sequence has been trimmed to remove vector sequence
and contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders Also
see Close TJ, Wing R, Kleinbols A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/gpages/bgn/31/cover.html)"
BASE COUNT 146 a 206 c 226 g 126 t
ORIGIN

Query Match 2.9%; Score 39.2; DB 10; Length 704;
Best Local Similarity 50.0%; Pred. No. 4.9;
Matches 98; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

QY 908 cggatcgaacgcgcgacatacacaagcgacgattatttgagcgtacacacatc 967
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 164 CGGTGATCTTCACGCGGACGACGCGCGCGTTCGCGTTCGACACAGA 223

QY 968 agattccgtatcgaagaagccgcgacgaagcctcgttcgctggttcgcgcgcagc 1027
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 224 AGATGAGCTCTGCTGAGCTGCGACGACATCGAGGCTTCGCGAAGTACGAGCCGGGC 283
QY 1028 cggacaatactcctacacgctacgacccctcgcgcacatttcggaanaaactcttca 1087
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 284 TGACCTGCTGACCGCGGACGCGGAGAGGCGCCAGATCATTACACACCTTGTGAGAGA 343
QY 1088 agttcagcaagccgt 1103
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 344 AGGCCAAGAGCGCGCT 359

RESULT 10
BF259476
LOCUS 759 bp mRNA linear EST 22-OCT-2001
DEFINITION HVSMEf0019D11f Hordeum vulgare seedling root EST library HVCNMA0007
(Etiolated and unstressed) Hordeum vulgare cDNA clone
HVSMEf0019D11f, mRNA sequence.
ACCESSION BF259476 GI:13119939
VERSION
KEYWORDS
SOURCE
ORGANISM
Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae
EST.
Hordeum vulgare
barley.
REFERENCE 1 (bases 1 to 759)
Wing, R., Close, T.J., Kleinbols, A., Wise, R., Begum, D., Frisch, D., Yu
, Y., Henry, D., Palmer, M., Rambo, T., Simmons, J., Choi, D.W., Fenton
, R.D., Oates, R. and Main, D.
Development of a genetically and physically anchored EST resource
for barley genomics: Morex unstressed seedling root cDNA library
Unpublished (2001)
JOURNAL On Nov 16, 2000 this sequence version replaced gi:1188505.
COMMENT Contact: Wing RA
Clemson University Genomics Institute
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 469
Seq primer: AATTAACCTCCTCAAGG
High quality sequence stop: 663.
FEATURES
Source
1..759
/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVSMEf0019D11f"
/clone_lib="Hordeum vulgare seedling root EST library
HVCNMA0007 (Etiolated and unstressed)"
/tissue_type="Seedling root"
/lab_host="TJC121"
/note="Vector: lambdaZAP; Site 1: EcoRI; Site 2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedling roots were
purified, total RNA was prepared, poly(A) RNA was
then harvested, total RNA was prepared, poly(A) RNA was
purified, one primary unamplified cDNA library was made,
and 1 million pfu were in vivo excised to give plusescript
SK(-) cDNA phagemids. These steps were performed in the TJ
Close laboratory at the University of California,
Riverside (Choi, Close, Fenton). Phagemids were plated and
picked at the Clemson University Genomics Institute (CUGI)
(Begum, Palmer, Frisch, Atkins and Wing). Plasmid DNA
preparations, DNA sequencing and sequence analysis were
performed at CUGI (Wing, Yu, Frisch, Henry, Simmons, Oates
, Rambo, Main). The sequence has been trimmed to remove
vector sequence and contains a minimum of 100 bases of
phred value 20 or above. For more details on library

Query Match	2.9%	Score 39.2	DB 10	Length 864
Best Local Similarity	50.0%	Pred. No. 5.4		
Matches	98	Conservative	0	Mismatches 98
				Indels 0
				Gaps 0
QY	908	cggataltgaacgycgcgcatattacacaaggcgcgcacgattatttggacgcgtacacatc	967	
Db	114	CGGTGATCTTTCAGCGCCGAGACACGCGGCACGCGGTTCACGTCTTCGGTTTGACAAAG	173	
QY	968	aggatttcogtltacgaagaagcgcgcacgaagaagctgttcgcttgggttgcgcagc	1027	
Db	174	AGATTGAGACTCTGTGACGCTGGCGGACGACATCGAGGCTTTCGCCAAGTGGACCCGGCG	233	
QY	1028	cggacaatactactcaatcaacgcgtlaagacccttcggcatttccttgaanaaacaacttca	1087	
Db	234	TGAGCTCGTACCGCGGACCGCGGACGAGGAGCGCGCAAGTCAATACACCCCTCTGGAGA	293	
QY	1088	agttcagcagaagccgt	1103	
Db	294	AGGCCAAGAGCGCCGT	309	

	RESULT	16
AM983120	LOCUS	
	DEFINITION	AM983120 888 bp mRNA linear EST 22-OCT-2001
	ACCESSION	HVSM60008D22f Hordeum vulgare pre-anthesis spike EST library
	VERSION	HVCDDNA0008 (white to yellow anther) Hordeum vulgare cDNA clone
	KEYWORDS	HVSM60008D22f, mRNA sequence.
	SOURCE	AM983120.3 GI:16317593
ORGANISM		EST.
REFERENCE		Barley.
AUTHORS		Hordeum vulgare
TITLE		Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
JOURNAL		Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
COMMENT		1 (bases 1 to 888).
		'T., Henry,D., Palmer,M., Rambo,T., Simmons,J., Chol,D.W., Fenton
		,R.D., Close,S.J., Oates,R. and Main,D.
		Development of a genetically and physically anchored EST resource
		for barley genomics: Morex pre-anthesis spike cDNA library
		Unpublished (2001)
		On Jun 2, 2000 this sequence version replaced gi:1153658.
		Contact: Wing RA
		Clemson University Genomics Institute
		Clemson University

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/organism="Hordeum vulgare"
/cultivar="Morex"
/db_xref="taxon:4513"
/clone="HVMEG0008D22f"
/clone_lib="Hordeum vulgare pre-anthesis spike EST library
HVCNA0008 (white to yellow anther)"
/tissue_type="pre-anthesis spike"
/lab_host="SOLR"
/notes="Vector: lambdaZAP, Site_1: EcoRI, Site_2: XhoI,

```

primary unamplified cDNA library was made, and 1 million pfu were in vivo excised to give Bluescript SK(-) cDNA Phagemids. These steps were performed in the TJ Close Lab (Choi) at the university of California, Riverside. Phagemids were plated and picked at the Clemson University Genomics Institute (CUGI) (Begum, Palmer, Frisch, Atkins and Wing) Plasmid DNA preparations, DNA sequencing and sequence analysis were performed at CUGI (Wing, Yu, Frisch , Henry, Simmons, Oates, Rambo, Main). The sequence has been trimmed to remove vector sequence and contains a minimum of 100 bases of phred value 20 or above. For more details on library preparation and sequence analysis see <http://www.genome.clemson.edu/projects/barley>. To order this clone see <http://www.genome.clemson.edu/orders> Also see Close TJ, Wing R, Kleinmorts A, Wise R (2001) Genetically and physically anchored EST resources for barley genomics. Barley Genetics Newsletter 31:29-30. (<http://wheat.pw.usda.gov/g9pages/bgn/31/cover.html>)"

Query Match	2.9%	Score 39.2	DB 9	Length 888
Best Local Similarity	50.0%	Pred.No.5.5		
Matches 98	Conservative 0	Mismatches 98	Indels 0	Gaps 0
QY 908	cggatttgaagcgcgcgattacacaagaagcgcgcacagattatttgagagcctaccacaac	967		
Db 166	CGGTGATCTTCGACGCCGGAACAGCGGGGACGCGGTGCACCTCTTCGGTTCGACAA	225		
QY 968	agatttcggtatcgaagaagaagcgcgaagaagctgttcggtggtgttcgcgcgaagc	1027		
Db 226	AGATGAGAGCTCTCCACGCTCGCGCGACGACATGAGAGTCTTCCCAAGGAGGAGCGCGGC	285		
QY 1028	cggaaacaaactcactcaacgcgtacagcccttggccatttctctgtaaaaaaactctta	1087		
Db 286	TGAGTCTGATCCCGCGAGCGCGCGAGAGCGCCGACAGTCCATCACACCCCTGTGAGA	345		
QY 1088	agttcacgacagccgt 1103			
Db 346	AGGCCAAGACGCCCT 361			
RESULT 17				
AI399239				
LOCUS	AI399239	506 bp	mRNA	linear
DEFINITION	NCM10F2T3 Westergaards Neurospora crassa	CDNA clone	W10F2 5'	mRNA
ACCESSION	AI399239			


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High quality sequence stop: 681
FEATURES
Location/Qualifiers
source
1. .697

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Db 129 CGCCGACGCGCCGAGAGCGCCGACATCATCACCCCTGTGTGAGAGGCCAAG 188
 QY 1098 agccgt 1103
 Db 189 CGCGGT 194

RESULT 25
 AZ049555/c 430 bp DNA linear GSS 06-MAR-2001
 LOCUS GSSBrn01308 Sheared genomic library Brucella melitensis biovar
 DEFINITION Abortus genomic clone M26, DNA sequence.
 ACCESSION AZ049555
 VERSION AZ049555.1 GI:7273470
 KEYWORDS GSS.
 SOURCE Brucella melitensis biovar Abortus.
 ORGANISM Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group; Brucellaceae; Brucella.

REFERENCE
 AUTHORS Sanchez D.O., Zandomeni R.O., Cravero S., Verdun R.E., Pierou E., Faccio P., Diaz G., Lanzavecchia S., Aguero F., Frasch A.C.C., Andersson S.G.E., Rosetti O.L., Grau O., and Ugalde R.A.
 TITLE Gene discovery through genomic sequencing of Brucella abortus
 JOURNAL Infect. Immun. 69 (2), 865-868 (2001)
 COMMENT Centro de Investigacion en Ciencias Agropecuarias (CICA) Instituto Nacional de Tecnologia Agropecuaria (INTA) C.C. 25 (1712) Castelar, Buenos Aires, Argentina
 Tel: 5411-4621-3316/1683
 Fax: 5411-4481-1316
 Email: zandomeni@inta.gov.ar
 Class: shotgun.

FEATURES
 source Location/Qualifiers
 1..430
 /organism="Brucella melitensis biovar Abortus"
 /strain="S-2308"
 /db_xref="taxon:235"
 /clone="M26"
 /clone_lib="Sheared genomic library"
 /note="Vector: Bluescript SK(-) (STRATAGENE); Genomic DNA was mechanically sheared, blunt ended, and size-fractionated by agarose gel electrophoresis. Fragments between 1.5-3 Kb were recovered and ligated to the EcoRV site of the Bluescript SK (-) vector."

BASE COUNT 83 a 124 c 127 g 91 t 5 others
 ORIGIN

Query Match 2.8%; Score 37; DB 12; Length 430;
 Best Local Similarity 51.3%; Pred. No. 16;
 Matches 79; Conservative 0; Mismatches 75; Indels 0; Gaps 0;

QY 442 gccatcttcgtcatgcatgacccaatccgtctgagcagaccctgtgtgtatc 501
 Db 221 gccatcttcgtcatgcatgacccaatccgtctgagcagaccctgtgtgtatc 162
 QY 502 aaagaagccgncgancatttcagcagangtctgtgcatgagccgtttaccgagcgt 561
 Db 161 tttagatccgcccgcgaatttcagtcagacccttcacgcatcagttcggttaagccgagc 102
 QY 562 aaatccatgtgtgtaaggcagctggcgagagc 595
 Db 101 aatattccgcaaacgacgacactgcccacccg 68

RESULT 26
 P947R 497 bp DNA linear GSS 25-JUL-2000
 LOCUS Leishmania major Friedlin PAC P947 right end-sequence, genomic
 DEFINITION survey sequence.
 ACCESSION AL390645
 VERSION AL390645.1 GI:9501621

KEYWORDS GSS.
 SOURCE Leishmania major.
 ORGANISM Eukaryota; Eukaryota; Kinetoplastida; Trypanosomatidae; Leishmania.

REFERENCE
 AUTHORS Ivens A.C., Lewis S.M., Bagherzadeh A., Zhang L., Chan H.M. and Smith D.F.
 TITLE A physical map of the Leishmania major Friedlin genome
 JOURNAL Genome Res. 8 (2), 135-145 (1998)
 MEDLINE 98146435
 REFERENCE 2 (bases 1 to 497)
 AUTHORS Taylor R.G., Huckle E.E.J., Ivens A.C., Rejandream M.A. and Barrell B.G.
 TITLE Direct Submission
 JOURNAL Submitted (24-JUL-2000) Leishmania major Friedlin genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and alicates@sanger.ac.uk
 see http://www.edi.ac.uk/parasites/leish.html
 details of Leishmania sequencing at the Sanger Centre are available at http://www.sanger.ac.uk/Projects/Lmajor/
 The primer sequence can be obtained from alicates@sanger.ac.uk.

COMMENT

FEATURES
 source Location/Qualifiers
 1..497
 /organism="Leishmania major"
 /strain="Friedlin"
 /db_xref="taxon:5664"
 /clone="PAC P947"

BASE COUNT 74 a 172 c 145 g 104 t 2 others
 ORIGIN

Query Match 2.8%; Score 37; DB 12; Length 497;
 Best Local Similarity 58.7%; Pred. No. 17;
 Matches 64; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

QY 1172 ctacctgtttgcgcgattatcgttcgcatcacgacgagcgagcagattgggtt 1231
 Db 231 CGACGCTGCTGGCGCACGGGCTGCGCGCCGCGCCGCGCGCTGAGCGCC 290
 QY 1232 gcttgaattgacgaagaagaccctgcttgcagcttcgttcgccc 1280
 Db 291 GCCAGCAACAGACGACACCCCGCTGTGCATGTCGTCGTCGTCCTCC 339

RESULT 27
 AL523270 893 bp mRNA linear EST 13-FEB-2001
 LOCUS AL523270 LTL_NFL003_NMC3 Homo sapiens cDNA clone CS0DC001YH12 5
 DEFINITION prime, mRNA sequence.
 ACCESSION AL523270
 VERSION AL523270.1 GI:12786763
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE
 AUTHORS Li W.B., Gruber C., Jessee J. and Polayes D.
 TITLE Full-length cDNA libraries and normalization
 JOURNAL Unpublished (2001)
 COMMENT Contact: Genoscope
 Genoscope - Centre National de Sequencage
 BP 191 91006 Evry cedex - France
 Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.

FEATURES
 source Location/Qualifiers
 1..893
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="CS0DC001YH12"
 /clone_lib="LTL_NFL003_NMC3"
 /sex="male"

/tissue_type="neuroblastoma cells"
/lab_host="DH10B"
/note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a Nott-oligo(dT) primer. Five prime end enriched, double-stranded cDNA was digested with Not I and cloned into the Not I and Eco RV sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies. Contact: Peng Liang Life Technologies, a division of Invitrogen, 9800 Medical Center Drive Rockville, Maryland 20850, USA Fax: (1) 301 610 8371 Email: filiang@lifestech.com URL: http://fulllength.invitrogen.com"

BASE COUNT 168 a 282 c 153 t 5 others
ORIGIN

Query Match 2.7%; Score 36.8; DB 9; Length 893;
Best Local Similarity 54.0%; Pred. No. 26;
Matches 68; Conservative 2; Mismatches 56; Indels 0; Gaps 0;

QY 1215 cgcgaagcattggttgatgagcgaagaagacccttggtgagcttgc 1274
DB 258 CGGGGGTGCACACCGCCAGCCGCGCGGAGTGGGCTGCTGCTGCC 317
QY 1275 ctgccggggaacatgacgaatgagccgctgtgctgaagtgctggaacnttgagaa 1334
DB 318 CAACCGCGGCACATTCGAGGAGTGCACCGAAGTCAAGAGAGCTTCCATTCAGAT 377
QY 1335 ggaag 1340
DB 378 GGAAGG 383

RESULT 28
AO161443/c 612 bp DNA linear GSS 09-SEP-1998
LOCUS mgx0007D20r CUGI Rice Blast BAC Library Magnaporthe grisea genomic
DEFINITION clone mgx0007D20r, DNA sequence.
ACCESSION AO161443
VERSION AO161443.1 GI:3557844
KEYWORDS GSS.
SOURCE Magnaporthe grisea.
ORGANISM Magnaporthe grisea.
Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes; Sordariomycetes incertae sedis; Magnaportheaceae; Magnaporthe.

REFERENCE 1 (bases 1 to 612)
Yu, Y., Zhu, H., Boyd, C.A., Gaudette, B., Gayle, A., Kingsbury, R., Phillips, K., Sasinowski, M., Wing, R.A. and Dean, R.A.

A BAC End Sequencing Framework to Sequence the Magnaporthe grisea Genome
Unpublished (1998)
JOURNAL Contact: Dean RA
COMMENT Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson University, Clemson, SC 29634
Tel: 864 656 5737
Fax: 864 656 4293
Email: rdean@clemson.edu
Seq primer: GGAACAGCTATGACCATG
Class: BAC ends
High quality sequence stop: 258.

FEATURES

source
1..612
Location/Qualifiers
/organism="Magnaporthe grisea"
/strain="70-15"
/db_xref="taxon:148305"
/clone="mgx0007D20r"
/clone_lib="CUGI Rice Blast BAC Library"
/tissue_type="Protoplasts"
/lab_host="E. coli DH10B"
/note="Vector: pBACWICH; Site_1: HindIII; Site_2: HindIII; Rice blast is one of the most devastating fungal diseases of rice world wide. It is a filamentous ascomycete with

a haploid genome (n=7) of approximately 40 Mbp. Rice blast is an important model fungal pathogen for studying numerous aspects of the fungal-host interaction. In order to facilitate genome wide analysis, a BAC library containing 9216 clones with an average insert size of 130 kbp was constructed. This library represents greater than 25x genome coverage. High density colony filters are available upon request."

BASE COUNT 118 a 233 c 124 g 137 t
ORIGIN

Query Match 2.7%; Score 36.6; DB 12; Length 612;
Best Local Similarity 44.8%; Pred. No. 25;
Matches 120; Conservative 0; Mismatches 148; Indels 0; Gaps 0;

QY 138 ggaagcgatgcgttcaaaaaaggccaagtgctgttgaagacaaaagatccggcgct 197
DB 589 GGTGGGCAATGTCGTTGACACGCTGTGATGCTGCTGAACCTCGGCTGTGAGGGGTGG 530
QY 198 ggtgttccgcgcgcngtltcaggaacaaatccgcgcacatccatcgcggaagcgct 257
DB 529 CGGCGCCACCGTCTGTGGGTGTACACGATACCGGACACGACGACGCGCGCAAGGCAC 470
QY 258 acttaagtcggtcgtgattgctgcttgaaagcaagcaatcgagttcgagcgtacgc 317
DB 469 CGGCATTCACCCGACGATGTTGACGACGACGCGCTTGCGCGGCTCGGACGACGATGAGA 410
QY 318 gccgaagcgttgcgaactaagcgcgcgaangaaatngnngcaatctgacatccatcg 377
DB 409 AGCGCGCGCCAGCATGTGTCGGCGGCGGAGCGCGCGATGATGACGCGGCGCAACCGG 350
QY 378 ttgtgactgcgtctgctgancgtcg 405
DB 349 TCGGGTCTGTGCGCGGACGACCGCG 322

RESULT 29
BM375129 388 bp mRNA linear EST 10-JAN-2002
LOCUS Ebem06_S0002_H03_R IGF barley Ebem06 library Hordeum vulgare cDNA
DEFINITION clone Ebem06_S0002_H03 5', mRNA sequence.
ACCESSION BM375129
VERSION BM375129.1 GI:18118519
KEYWORDS EST.
SOURCE barley.
ORGANISM Hordeum vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae ; Triticeae; Hordeum.

REFERENCE 1 (bases 1 to 388)
Hedley, P., Liu, H., Caldwell, D., McCallum, N., Mudie, S., Cardle, L., Ramsay, L., Machray, G., Marshall, D.F.M. and Waugh, R.
Development of Barley Transcriptome Resources
Unpublished (2001)
Contact: Waugh R
Unit of Genomics
Scottish Crop Research Institute
Invergowrie, Dundee, DD2 5DA, Scotland, UK
Tel: 00 44 1382 562731
Fax: 00 44 1382 562426
Email: rwaugh@scri.sari.ac.uk
All sequence has a phred quality score of 20 or over
Seq primer: M13 reverse.

FEATURES

source
1..388
Location/Qualifiers
/organism="Hordeum vulgare"
/cultivar="Optic"
/db_xref="taxon:4513"
/clone="Ebem06_S0002_H03"
/clone_lib="IGF barley Ebem06 library"
/tissue_type="Embryo"
/dev_stage="21 days post anthesis"

Query Match 2.7%; Score 36; DB 9; Length 488;
Best Local Similarity 58.3%; Pred. No. 33;
Matches 63; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

Qy 608 atgtcgcaacatcgaaacatgaattcgagcgccgcatccggcgtttgagtga 667
Db 222 ATGCTACCCGCTCCCTCCATCTGTGTGCGGACCCCGTGAAGAGAGAGCTTGA 281

Qy 668 cgcacatcattcattgagccggtcggtgcaacaacacggtttga 715
Db 282 CGCCCAAGTGCCTGCAATTCGAGCTCGGAAGAAGCAAGCCCACTTGA 329

RESULT 41
AT101002 497 bp mRNA linear EST 18-DEC-1999
LOCUS we09e08.x1 NCI-CGAP_Lu24 Homo sapiens cDNA clone IMAGE:2340614.3
DEFINITION similar to TR:013863 013863 HYPOTHETICAL 60.7 KD PROTEIN C1B1.02C
IN CHROMOSOME 1.; contains element MER22 MER22 repetitive element
; mRNA sequence.

ACCESSION AT101002 GI:4988902
VERSION AT101002.1
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 497)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bdrp/image/image.html
Insert Length: 760 Std Error: 0.00
Seq primer: -400p from Gibco
High quality sequence stop: 461.
Location/Qualifiers

FEATURES
source
1. 497
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2340614"
/clone_lib="NCI-CGAP_Lu24"
/tissue_type="carcinoid"
/lab_host="DH10B"
/note="Organ: lung; Vector: pRT73D-Pac (Pharmacia) with a
modified polylinker; Plasmid DNA from the normalized
library NCI-CGAP_Lu5 was prepared, and ss circles were
made in vitro. Following HAP purification, this DNA was
used as tracer in a subtractive hybridization reaction.
The driver was PCR-amplified cDNAs from a pool of 5,000
clones made from the same library (clonoids
1414920-1417991 and 1520904-1522439). Subtraction by Bento
Soares and M. Fatima Bonaldo.

BASE COUNT 98 a 162 c 144 g 93 t
ORIGIN

Query Match 2.7%; Score 36; DB 9; Length 497;
Best Local Similarity 58.3%; Pred. No. 33;
Matches 63; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

Qy 608 atgtcgcaacatcgaaacatgaattcgagcgccgcatccggcgtttgagtga 667
Db 222 ATGCTACCCGCTCCCTCCATCTGTGTGCGGACCCCGTGAAGAGAGAGCTTGA 281

Qy 668 cgcacatcattcattgagccggtcggtgcaacaacacggtttga 715
Db 282 CGCCCAAGTGCCTGCAATTCGAGCTCGGAAGAAGCAAGCCCACTTGA 329

RESULT 42
AM006812 526 bp mRNA linear EST 10-SEP-1999
LOCUS w070712.x1 NCI-CGAP_C03 Homo sapiens cDNA clone IMAGE:2506799.3
DEFINITION similar to TR:013863 013863 HYPOTHETICAL 60.7 KD PROTEIN C1B1.02C
IN CHROMOSOME 1.; contains element MER22 MER22 repetitive element
; mRNA sequence.

ACCESSION AM006812 GI:5855590
VERSION AM006812.1
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 526)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D.,
Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bdrp/image/image.html
Seq primer: -400p from Gibco
High quality sequence stop: 470.
Location/Qualifiers

FEATURES
source
1. 526
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2506799"
/clone_lib="NCI-CGAP_C03"
/sex="pooled"
/tissue_type="colon"
/lab_host="DH10B"
/note="Vector: pRT73D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from 12 pooled bulk tumor samples and primed
with a Not I - oligo(dT) primer. Double-stranded cDNA was
ligated to Eco RI adaptors (Pharmacia), digested with Not
I and cloned into the Not I and Eco RI sites of the
modified pRT73 vector. Library went through one round of
normalization."

BASE COUNT 106 a 165 c 143 g 111 t 1 others
ORIGIN

Query Match 2.7%; Score 36; DB 9; Length 526;
Best Local Similarity 58.3%; Pred. No. 34;
Matches 63; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

Qy 608 atgtcgcaacatcgaaacatgaattcgagcgccgcatccggcgtttgagtga 667
Db 204 ATGCTACCCGCTCCCTCCATCTGTGTGCGGACCCCGTGAAGAGAGAGCTTGA 263

Qy 668 cgcacatcattcattgagccggtcggtgcaacaacacggtttga 715
Db 264 CGCCCAAGTGCCTGCAATTCGAGCTCGGAAGAAGCAAGCCCACTTGA 311

RESULT 43
AT1743812

